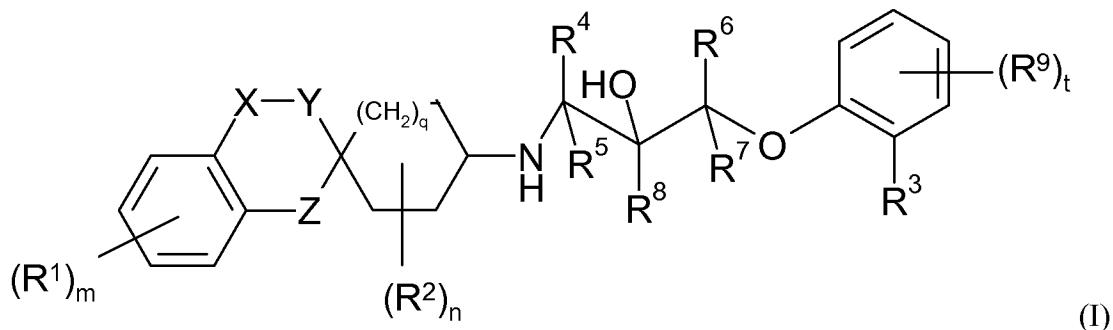


Amendments to the Claims:

This listing of claims replaces all prior versions and listings of claims in the application:

Listing of Claims:

1. (Currently amended) A compound of formula



wherein

m is 0, 1, 2, 3 or 4;

each R¹ independently represents halogen, cyano, hydroxyl, C₁-C₆ alkyl, C₁-C₆ haloalkyl, C₁-C₆ alkoxy or sulphonamido; either X represents a bond, -CH₂-, -O- or -C(O)- and Y represents a bond, -CH₂-, -O- or -C(O)-, or X and Y together represent a group -CH=C(CH₃)₂ or -C(CH₃)=CH-, and Z represents a bond, -O-, -NH- or -CH₂-; provided that only one of X, Y and Z can represent a bond at any one time and provided that X and Y do not both simultaneously represent -O- or -C(O)-;

n is 0, 1 or 2;

each R² independently represents halogen or C₁-C₆ alkyl;

q is 0 or 1;

R³ represents -NHC(O)R¹⁰, -C(O)NR¹¹R¹² or -COOR^{12a};

R^4 , R^5 , R^6 , R^7 and R^8 each independently represent a hydrogen atom or a C_1-C_6 -alkyl group;

t is 0, 1 or 2;

each R^9 independently represents halogen, cyano, hydroxyl, carboxyl, C_1-C_6 alkoxy, C_1-C_6 alkoxy carbonyl, C_1-C_6 haloalkyl, or C_1-C_6 alkyl optionally substituted by at least one substituent selected from carboxyl and C_1-C_6 alkoxy carbonyl;

R^{10} represents a group C_1-C_6 alkyl, C_2-C_6 alkenyl, C_3-C_6 cycloalkyl, adamantlyl, C_5-C_6 cycloalkenyl, phenyl or a saturated or unsaturated 5- to 10-membered heterocyclic ring system comprising at least one ring heteroatom selected from nitrogen, oxygen and sulphur, each of which may be optionally substituted by one or more substituents independently selected from nitro, hydroxyl, oxo, halogen, carboxyl, C_1-C_6 alkyl, C_1-C_6 alkoxy, C_1-C_6 alkylthio, C_1-C_6 alkyl carbonyl, C_1-C_6 alkoxy carbonyl, phenyl and $-NHC(O)-R^{13}$, or

R^{10} represents a group $-NR^{14}R^{15}$ or $-O-R^{16}$;

R^{11} and R^{12} each independently represent (i) a hydrogen atom, (ii) a 3- to 6-membered saturated or unsaturated ring optionally comprising at least one ring heteroatom selected from nitrogen, oxygen and sulphur and optionally further comprising a bridging group, the ring being optionally substituted with at least one substituent selected from halogen, hydroxyl, C_1-C_6 alkyl, C_1-C_6 hydroxy alkyl and C_1-C_6 haloalkyl,

(iii) a C_1-C_6 alkyl group optionally substituted by at least one substituent selected from halogen, amino, hydroxyl, C_1-C_6 haloalkyl, carboxyl, C_1-C_6 alkoxy, C_1-C_6 alkoxy carbonyl, C_1-C_6 alkyl carbonyl amino and a 3- to 6-membered saturated or unsaturated ring optionally comprising at least one ring heteroatom selected from nitrogen, oxygen and sulphur and optionally further comprising a bridging group, the ring being optionally substituted with at least one substituent selected from halogen, hydroxyl, oxo, C_1-C_6 alkyl, C_1-C_6 hydroxy alkyl and C_1-C_6 haloalkyl, or (iv) C_1-C_6 alkylsulphonyl,

or

R^{11} and R^{12} together with the nitrogen atom to which they are attached form a 4- to 7-membered saturated heterocyclic ring that optionally further comprises a ring nitrogen, oxygen or sulphur atom and that is optionally fused to a benzene ring to form a 8- to 11-membered ring system, the heterocyclic ring or ring system being optionally substituted with at least one substituent selected from halogen, hydroxyl, amido, C_1 - C_6 alkyl, C_1 - C_6 hydroxyalkyl, C_1 - C_6 alkoxy, C_1 - C_6 alkoxy carbonyl, C_1 - C_6 haloalkyl, C_1 - C_6 alkylamino, di- C_1 - C_6 alkylamino, C_1 - C_6 alkylcarbonyl, C_1 - C_6 alkylcarbonyl amino, C_1 - C_6 alkylaminocarbonyl, di- C_1 - C_6 alkylaminocarbonyl, phenyl, halophenyl, phenylcarbonyl, phenylcarbonyloxy and hydroxydiphenylmethyl;

R^{12a} represents a hydrogen atom or a C_1 - C_6 alkyl group;

R^{13} represents a C_1 - C_6 alkyl, amino or phenyl group;

R^{14} and R^{15} each independently represent a hydrogen atom, or a group C_1 - C_6 alkyl, C_1 - C_6 alkylsulphonyl, phenyl or a saturated or unsaturated 5- to 10-membered heterocyclic ring system comprising at least one ring heteroatom selected from nitrogen, oxygen and sulphur, each group being optionally substituted as defined above for R^{10} , or

R^{14} and R^{15} together with the nitrogen atom to which they are attached form a 4- to 7-membered saturated heterocyclic ring that optionally further comprises a ring nitrogen, oxygen or sulphur atom, the heterocyclic ring being optionally substituted by at least one hydroxyl; and

R^{16} represents a hydrogen atom, or a group C_1 - C_6 alkyl, phenyl or a saturated or unsaturated 5- to 10-membered heterocyclic ring system comprising at least one ring heteroatom selected from nitrogen, oxygen and sulphur, each group being optionally substituted as defined above for R^{10} ;

or a pharmaceutically acceptable salt or solvate thereof.

2. (Original) A compound according to claim 1, wherein X and Y have the meanings shown in the following table:

X	Y
bond	O
O	bond
CH ₂	bond
bond	CH ₂

3. (Previously presented) A compound according to claim 1, wherein Z represents -O- or -CH₂-.

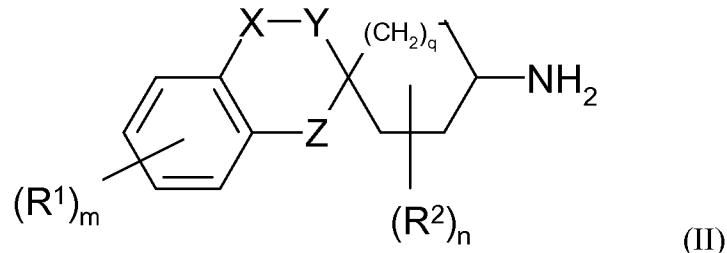
4. (Previously presented) A compound according to claim 1, wherein q is 1.

5. (Previously presented) A compound according to claim 1, wherein R³ represents -NHC(O)R¹⁰ or -C(O)NR¹¹R¹².

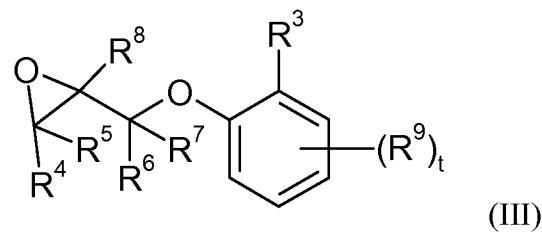
6. (Previously presented) A compound according to claim 1, wherein t is 1 and R⁹ is located in the *para* position with respect to R³.

7. (Original) A compound according to claim 1 selected from:
2-((2S)-3-[(5-Chloro-3H-spiro[1-benzofuran-2,1'-cyclohexan]-4'-yl)amino]-2-hydroxypropyl}oxy)-4-hydroxy-N-methylbenzamide,
N-2-((2S)-3-[(5-Chloro-3H-spiro[1-benzofuran-2,1'-cyclohexan]-4'-yl)amino]-2-hydroxypropyl}oxy)-4-fluorophenyl]acetamide,
2-((2S)-3-[(5-Chloro-3H-spiro[1-benzofuran-2,1'-cyclohexan]-4'-yl)amino]-2-hydroxypropyl}oxy)-N-methylbenzamide,
N-[2-((2S)-3-[(5-Chloro-3H-spiro[1-benzofuran-2,1'-cyclohexan]-4'-yl)amino]-2-hydroxypropyl}oxy)-4-hydroxyphenyl]acetamide,
N-[2-((2S)-3-[(5-Chloro-3H-spiro[1-benzofuran-2,1'-cyclohexan]-4'-yl)amino]-2-hydroxy-2-methylpropyl}oxy)-4-hydroxyphenyl]acetamide (trifluoro acetate),
and pharmaceutically acceptable salts and solvates of any one thereof.

8. (Original) A process for the preparation of a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof as defined in claim 1 which comprises,
(a) reacting a compound of formula

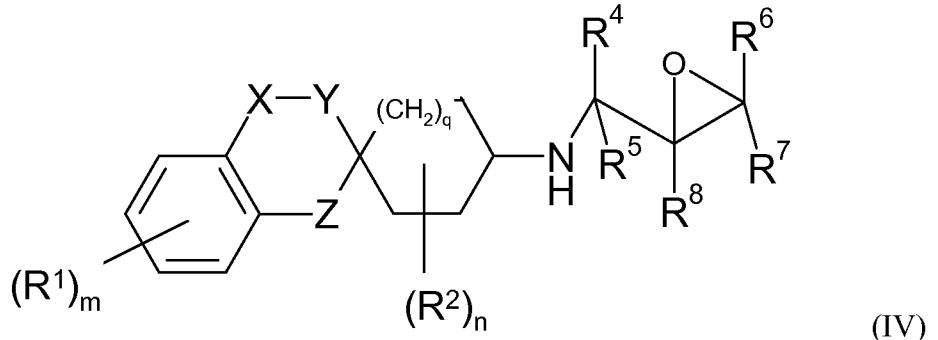


wherein m, R¹, n, R², q, X, Y and Z are as defined in formula (I), with a compound of formula

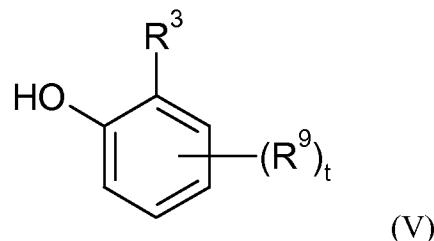


wherein R³, R⁴, R⁵, R⁶, R⁷, R⁸, t and R⁹ are as defined in formula (I); or

(b) reacting a compound of formula

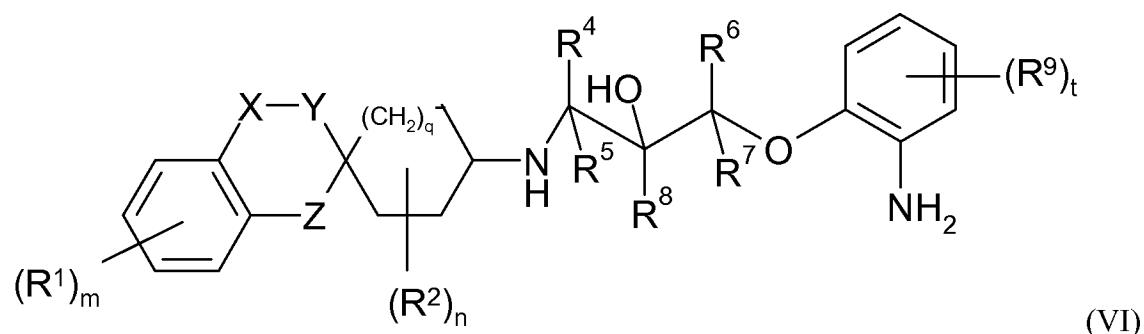


wherein m, R¹, n, R², q, X, Y, Z, R⁴, R⁵, R⁶, R⁷ and R⁸ are as defined in formula (I), with a compound of formula

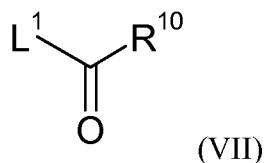


wherein R³, t and R⁹ are as defined in formula (I), in the presence of a suitable base; or

(c) when R³ represents -NHC(O)R¹⁰, reacting a compound of formula

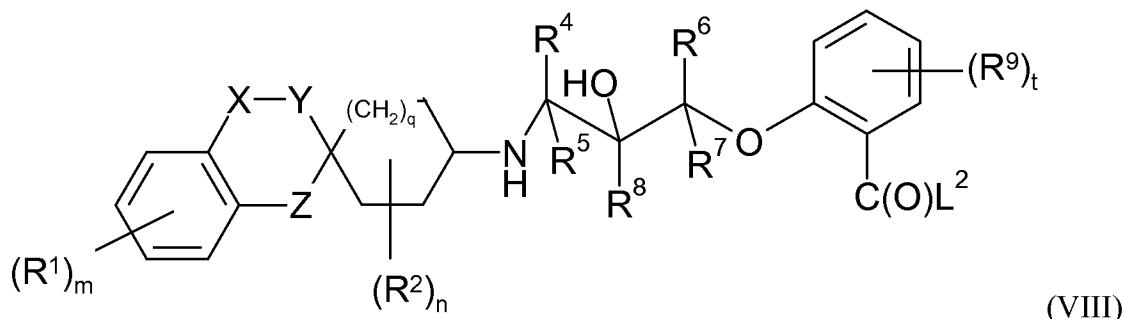


wherein m, R¹, n, R², q, X, Y, Z, R⁴, R⁵, R⁶, R⁷, R⁸, t and R⁹ are as defined in formula (I), with a compound of formula



wherein L¹ represents a leaving group and R¹⁰ is as defined in formula (I); or

(d) when R³ represents -C(O)NR¹¹R¹², reacting a compound of formula



wherein L² represents a leaving group and m, R¹, n, R², q, X, Y, Z, R⁴, R⁵, R⁶, R⁷, R⁸, t and R⁹ are as defined in formula (I), with a compound of formula (IX), NHR¹¹R¹², wherein R¹¹ and R¹² are as defined in formula (I); or

(e) when R^3 represents $-NHC(O)R^{10}$, R^{10} represents $-NR^{14}R^{15}$ and R^{14} and R^{15} both represent hydrogen, reacting a compound of formula (VI) as defined in (c) above with potassium cyanate;

and optionally after (a), (b), (c), (d) or (e) forming a pharmaceutically acceptable salt or solvate.

9. (Currently amended) A pharmaceutical composition comprising a compound of formula (I) or a pharmaceutically acceptable salt ~~or solvate~~ thereof as claimed in claim 1 in association with a pharmaceutically acceptable adjuvant, diluent or carrier.

10. (Previously presented) A process for the preparation of a pharmaceutical composition as claimed in claim 9 which comprises mixing a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof as claimed in claim 1 with a pharmaceutically acceptable adjuvant, diluent or carrier.

11. (Cancelled)

12. (Previously presented) A method of treating a disease or condition in which modulation of chemokine receptor activity is beneficial, the method comprising administering to a patient in need thereof a therapeutically effective amount of a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof as claimed in claim 1.

13. (Previously presented) A method of treating rheumatoid arthritis, the method comprising administering to a patient in need thereof a therapeutically effective amount of a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof as claimed in claim 1.

14. (Previously presented) A method of treating chronic obstructive pulmonary disease, the method comprising administering to a patient in need thereof a therapeutically effective amount of a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof as claimed in claim 1.

15. (Previously presented) A method of treating asthma, the method comprising administering to a patient in need thereof a therapeutically effective amount of a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof as claimed in claim 1.

16. (Previously presented) A method of treating multiple sclerosis, the method comprising administering a therapeutically effective amount of a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof as claimed in claim 1.

17. (Previously presented) A method of treating an inflammatory disease which comprises administering to a patient in need thereof a therapeutically effective amount of a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof as claimed in claim 1.

18. (Previously presented) A method of treating an airways disease which comprises administering to a patient in need thereof a therapeutically effective amount of a compound of formula (I) or a pharmaceutically acceptable salt or solvate thereof as claimed in claim 1.